Pharmacy Technician Course

INTRODUCTION

SECTION 1: History and Scope of the Pharmacy Technician

SECTION 2: Manufacturing Pharmaceuticals

SECTION 3: Rules and Regulations (PTCB 2.0)

SECTION 4: Controlled Substances (PTCB 2.4)

SECTION 5: Pharmacy Safety

SECTION 6: Drug Classifications (PTCB Pharmacology 1.1 – 1.6)

SECTION 7: Sterile and Non-Sterile Compounding (PTCB 3.0 – 3.7)

SECTION 8: Pharmacy Quality Assurance (PTCB 5.1-5.5)

SECTION 9 – Mathematics in Pharmacy (PTCB 3.6, 6.3)

SECTION 10: Inventory Management (PTCB 7.1-7.5)

SECTION 11: Pharmacy Billing and Reimbursement (PTCB 8.0)

SECTION 12: Information Systems (PTCB 9.0)

Activity Summary

• Activity Title: Pharmacy Technician Course
• Release date: 2018-06-01
• Expiration date: 2021-06-01
• Estimated time to complete activity: 8 hours
• This course is accessible with any web browser. We recommend recent versions of Google Chrome, Internet Explorer 9 and later, or Apple iPad.
• This course is jointly provided by Pacific Medical Training and Postgraduate Institute for Medicine (PIM). You may reach PIM at inquiries@pimed.com.

Target Audience

This activity has been designed to meet the educational needs of pharmacy technicians, physicians, physician assistants, nurse practitioners, registered nurses, pharmacists and dentists who endeavor to take the Pharmacy Technician Certification Board (PTCB) exam.
Educational Objectives

After completing this activity, the participant should be better able to:

• Describe the history and scope of the pharmacy technician.
• Summarize the manufacturing process of pharmaceuticals.
• Identify the difference between brand and generic medication.
• Explain the Food and Drug Administration’s role in the approval process of bringing medication to market.
• Discuss the federal, state, and/or local rules and regulations of the pharmaceutical industry.
• Categorize controlled substances based on their potential for addiction and abuse.
• Identify forms for storing, maintaining, and distributing controlled substances.
• Recall prescribing and filing rules and regulations for controlled substances.
• Demonstrate pharmacy safety, such as storage and handling of hazardous substances.
• Describe the purpose of restricted drug programs.
• Identify the specific job duties that are only done by a licensed pharmacist.
• Underline common drug interactions.
• Distinguish drug classifications according to their function.
• Express the purpose and methods of sterile and non-sterile compounding of medications.
• Explain the importance and various aspects of pharmacy quality assurance.
• Demonstrate a general understanding of mathematics in pharmacy.
• Employ pharmacy inventory management.
• Summarize pharmacy billing and reimbursement.
• Discuss the use of pharmacy information systems that are in use to manage prescription needs of patients.

Faculty

• Judith Haluka, EMT-Paramedic – State of Pennsylvania

Joint Accreditation Statement

In support of improving patient care, this activity has been planned and implemented by the Postgraduate Institute for Medicine and Pacific Medical Training. Postgraduate Institute for
Medicine is jointly accredited by the Accreditation Council for Continuing Medical Education (ACCME), the Accreditation Council for Pharmacy Education (ACPE), and the American Nurses Credentialing Center (ANCC), to provide continuing education for the healthcare team.

Continuing Pharmacy Education

Postgraduate Institute for Medicine designates this continuing education activity for 8.0 contact hour(s) (0.80 CEUs) of the Accreditation Council for Pharmacy Education.

Universal Activity Number - JA4008162-9999-18-141-H04-T

Type of Activity: Knowledge

Disclosure of Conflicts of Interest

Postgraduate Institute for Medicine (PIM) requires instructors, planners, managers and other individuals who are in a position to control the content of this activity to disclose any real or apparent conflict of interest (COI) they may have as related to the content of this activity. All identified COI are thoroughly vetted and resolved according to PIM policy. PIM is committed to providing its learners with high quality CME activities and related materials that promote improvements or quality in healthcare and not a specific proprietary business interest of a commercial interest.

The faculty reported the following financial relationships or relationships to products or devices they or their spouse/life partner have with commercial interests related to the content of this CME activity:
• Judith Haluka — Has no real or apparent conflicts of interest to report

The PIM planners and managers have nothing to disclose. The Pacific Medical Training planners and managers have nothing to disclose.

Method of Participation and Request for Credit

During the period 2018-06-01 through 2021-06-01 participants must read the learning objectives and faculty disclosures and study the educational activity.

Upon successfully completing the post-test with a score of 84% or better and the activity evaluation form, transcript information will be sent to the NABP CPE Monitor Service within 4 weeks.

Media

Internet

Disclosure of Unlabeled Use

This educational activity may contain discussion of published and/or investigational uses of agents that are not indicated by the FDA. The planners of this activity do not recommend the use of any agent outside of the labeled indications.

The opinions expressed in the educational activity are those of the faculty and do not necessarily represent the views of the planners. Please refer to the official prescribing information for each product for discussion of approved indications, contraindications, and warnings.

Disclaimer

Participants have an implied responsibility to use the newly acquired information to enhance patient outcomes and their own professional development. The information presented in this activity is not meant to serve as a guideline for patient management. Any procedures, medications, or other courses of diagnosis or treatment discussed or suggested in this activity should not be used by clinicians without evaluation of their patient's conditions and possible
contraindications and/or dangers in use, review of any applicable manufacturer's product information, and comparison with recommendations of other authorities.